

Photochemical CO₂ reduction in water with a carbon nitride modified with Co porphyrin molecular catalyst

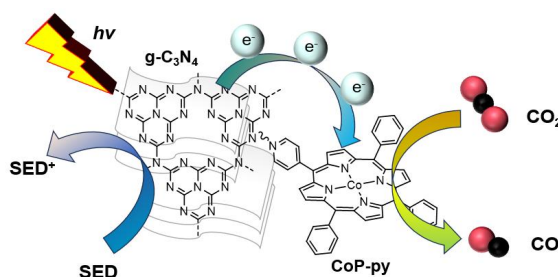
(¹Graduate School of Chemistry, Kyushu University)

○Guan Wang,¹ Sakai Ken,¹ Hironobu Ozawa¹

Keywords: CO₂ Reduction; Photocatalyst; Graphitic Carbon Nitride; Co porphyrin.

Recently, development of artificial photosynthetic systems, made up of a semiconductor photocatalyst and a functional molecule, promoting visible-light-driven CO₂ reduction reaction (CO₂RR) has received increasing attention. Among a large number of semiconductors, graphitic carbon nitride (g-C₃N₄) is often used as a semiconductor photocatalyst for CO₂RR because the conduction band edge potential possesses sufficiently negative potential for CO₂RR.¹ Nevertheless, a g-C₃N₄ photocatalyst modified with a molecular catalyst showing a high selectivity for CO₂ reduction even in a fully aqueous media is still very rare.

In this study, a g-C₃N₄ photocatalyst chemisorbed with Co porphyrin molecular catalyst having a pyridyl anchor² (g-C₃N₄/CoP-py) has been prepared and its performance for photocatalytic CO₂RR in the presence of a sacrificial electron doner (TEOA) has been evaluated in 0.1 M NaHCO₃ aqueous media saturated with CO₂. Under visible light irradiation conditions ($\lambda > 400$ nm), g-C₃N₄/CoP-py exhibits CO₂-to-CO conversion activity with a quite high CO production selectivity (about 80%, TON_{8h} = 3.4). This study revealed for the first time that our pyridyl anchoring technique is also effective for C₃N₄ and that higher selectivity is achieved by chemisorption of a relatively large amount of Co porphyrin promoting CO₂-to-CO conversion with a higher selectivity even in fully aqueous media³.



1) X. Wang, K. Maeda, A. Thomas, K. Takanabe, G. Xin, J. M. Carlsson, K. Domen, M. Antonietti, *Nat. Mater.*, **2009**, 8, 76-80.

2) H. Ozawa, R. Kikunaga, H. Suzuki, R. Abe, K. Sakai, *Sustain. Energy Fuels*, **2023**, 7, 1627-1632.

3) A. Call, M. Cibian, K. Yamamoto, T. Nakazono, K. Yamauchi, K. Sakai, *ACS Catal.* **2019**, 9, 4867-4874.